EMISSION MEASUREMENT CENTER TECHNICAL INFORMATION DOCUMENT

DERIVATION OF) H@

INTRODUCTION

The term $)H_{\scriptscriptstyle \odot}$ given in Equation 1 below is determined during the calibration of the orifice meter in the Environmental Protection Agency (EPA) Method 5 sampling train. The purpose of this document is to explain the derivation of this term.

$$) H_{@} = {}^{f} \frac{0.0319 \) H^{f} \left[(t_{w} + 460) \ 2 \right]^{2}}{P_{b} (t_{o} + 460) \left[V_{w} \right]^{f}}$$
 Eq. 1

DERIVATION

The flow rate through an orifice meter is defined by the following equation:

$$Q_{m} = K_{m} \frac{T_{m})H}{\sqrt{P_{m}M_{m}}}$$
 Eq. 2

where:

 Q_m = volumetric flow rate through the orifice meter, cfm.

 K_m = orifice meter calibration constant, [(in. Hg)(lb-mole)]/ [(°R)(in. H₂0)].

 T_m = absolute orifice meter temperature, °R.

) H^f = orifice meter pressure differential, in. H_2O .

 P_m = absolute orifice meter pressure, in. Hg.

 $M_{\rm m}$ = molecular weight of gas flowing through the orifice meter, lb/lb-mole.

Instead of using the orifice meter coefficient K_m , the term $)H_{\mbox{\tiny @}}$ was derived to reflect the general sampling conditions of sampling trains and orifice meter designs with a calibration constant that could be related to the desired flow rate. For example, a $)H_{\mbox{\tiny @}}$ of 1.84 means that the orifice pressure differential)H would be about 1.84 when sample gas is flowing through the orifice meter. The reference conditions for)H were chosen to be 0.75 dcfm of air at 68°F and 29.92 in. Hg. Using Equation 2, $)H_{\mbox{\tiny @}}$ is

then defined as:

)
$$H_{@} = \frac{Q_{@}2 P_{@} M_{@}}{K_{m}^{2} T_{@}}$$
 fEq. 3

where the subscript "@" represents the reference conditions of 0.75 dcfm of air at $68^{\circ}F$ and 29.92 in. Hg.

Substituting the values for the reference conditions (except $M_{\scriptscriptstyle \circledcirc}$) into Equation 3, the following is obtained:

)
$$H_{\odot} = \frac{(0.75^2) (29.92) M_{\odot}}{528 K_{m}^2} = 0.0319 \frac{M_{\odot}}{K_{m}^2}$$
 f Eq. 4

Now, the relationship between Q_m and the volumetric flow rate through the wet test meter (Q_w) is as follows (the subscript "w" refers to the flow rate conditions at the wet test meter):

$$\frac{P_m \ Q_m}{T_m} = \frac{P_w \ Q_w}{T_w}$$
 Eq. 5

Substituting Equation 2 into Equation 5 and solving for ${K_{\scriptscriptstyle m}}^2 \colon$

$$\frac{P_{m} K_{m}}{T_{m}} \frac{T_{m} H}{\int P_{m} M_{m}} = \frac{P_{w} Q_{w}}{T_{w}}$$

$$K_{m}^{2} = \frac{M_{m} T_{m} P_{w}^{2} Q_{w}^{2}}{) H T_{w}^{2} P_{m}}$$
 Eq. 6

In the Method 5 calibration set-up, the pressure at the orifice meter and the wet test meter are at barometric pressure. Thus, P_w = P_m = P_b , Equation 6 simplifies to:

$$K_{m}^{2} = \frac{M_{m} T_{m} P_{b} Q_{w}^{2}}{M_{m} T_{w}^{2}}$$
 Eq. 7

Substituting Equation 7 into Equation 4:

$$) H_{\odot} = \frac{0.0319 M_{\odot}) H T_{w}^{2}}{M_{m} T_{m} P_{b} O_{w}^{2}}$$
 ff Eq. 8

Since air is being used to calibrate the orifice, $M_{\odot}=M_{m}$. In addition, $Q_{w}=V_{w}/2$, which is the volume measured by the wet test meter divided by time, and T_{m} is the same temperature measured at the outlet of the dry gas meter, i.e., $T_{m}=T_{o}$. Furthermore, $T_{w}=t_{w}+460$, and $T_{o}=t_{o}+460$. Making these substitutions in Equation 8 yields:

where:

)H $_{\tiny \tiny \odot}$ =)H that gives 0.75 dcfm of air at 68°F and 29.92 in. Hg, in. H_2O .

 $0.0319^{f} = constant, (0.75^{2})(29.92)/(68 + 460).$

) H^f = orifice meter pressure differential, in. H_2O .

 t_w = wet test meter temperature, °F.

 2^{f} = time, min.

 P_b = barometric pressure, in. Hg.

t_o = outlet temperature of dry gas meter, °F.

 V_w = volume measured by the wet test meter, cf.

BIBLIOGRAPHY

- 1. Code of Federal Regulations. Title 40, Part 60. U.S. Government Printing Office, Washington, D.C. July 1, 1987.
- 2. Shigehara, R.T. Adjustments in the EPA Nomograph for Different Pitot Tube Coefficients and Dry Molecular Weights. In: Stack Sampling Technical Information A Collection of Monographs and Papers, Vol. III. U.S. Environmental Protection Agency. Publication No. EPA-450-2-78-042c. October 1978. pp. 48-49.
- 3. Smith, W.S., R.M. Martin, D.E. Durst, R.G. Hyland, T.J. Logan, and C.B. Hager. Stack Gas Sampling Improved and Simplified with New Equipment. Presented at the 60th Annual Meeting of the Air Pollution Control Association, June 11-16, 1967, Cleveland, Ohio. APCA Paper N. 67-119.